Attorney Docket No.: PB60565USw

## **Amendments To The Claims:**

This listing of claims will replace all prior versions, and listings, of claims in the application:

## What is claimed is:

1. (Original) A compound of formula (I):

$$R^{2a}$$
 $R^{2a}$ 
 $R^{8}$ 
 $R^{9}$ 
 $R^{9}$ 
 $R^{2a}$ 
 $R^{2a}$ 
 $R^{2a}$ 
 $R^{2a}$ 
 $R^{2a}$ 
 $R^{2a}$ 
 $R^{2a}$ 
 $R^{2a}$ 
 $R^{2a}$ 
 $R^{2a}$ 

wherein:

W represents N or CR<sup>10</sup> wherein R<sup>10</sup> represents hydrogen, halogen, optionally substituted alkyl, optionally substituted aryl, or optionally substituted heterocyclyl;

(1)

X represents N or CR<sup>11</sup> wherein R<sup>11</sup> represents hydrogen, halogen, optionally substituted alkyl, optionally substituted aryl, or optionally substituted heterocyclyl;

Y represents N or CR<sup>12</sup> wherein R<sup>12</sup> represents hydrogen, halogen, CH<sub>3</sub> or CF<sub>3</sub>;

Z represents O, S, SO or SO<sub>2</sub>;

R<sup>1</sup> represents CO<sub>2</sub>R<sup>4</sup>, CONR<sup>5</sup>R<sup>6</sup>, CH<sub>2</sub>CO<sub>2</sub>H, optionally substituted SO<sub>2</sub>alkyl, SO<sub>2</sub>NR<sup>5</sup>R<sup>6</sup>, NR<sup>5</sup>CONR<sup>5</sup>R<sup>6</sup>, 2*H*-tetrazol-5-yl-methyl or optionally substituted heterocyclyl;

R<sup>2a</sup> and R<sup>2b</sup> independently represents hydrogen, halo, optionally substituted alkyl, optionally substituted alkoxy, CN, SO<sub>2</sub>alkyl, SR<sup>5</sup>, NO<sub>2</sub>, optionally substituted aryl, CONR<sup>5</sup>R<sup>6</sup> or optionally substituted heteroaryl;

R<sup>x</sup> represents optionally substituted alkyl wherein 1 or 2 of the non-terminal carbon atoms are optionally substituted by a group independently selected from NR<sup>4</sup>, O and SO<sub>n</sub>, wherein n is 0, 1 or 2: or R<sup>x</sup> represents optionally substituted CQ<sup>a</sup>Q<sup>b</sup>-heterocyclyl, optionally substituted CQ<sup>a</sup>Q<sup>b</sup>-bicyclic heterocyclyl or optionally substituted CQ<sup>a</sup>Q<sup>b</sup>-aryl;

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R<sup>4</sup> represents hydrogen or an optionally substituted alkyl;

R<sup>5</sup> represents hydrogen or an optionally substituted alkyl;

R<sup>6</sup> represents hydrogen or optionally substituted alkyl, optionally substituted heteroaryl, optionally substituted SO<sub>2</sub>aryl, optionally substituted SO<sub>2</sub>alkyl, optionally substituted SO<sub>2</sub>heteroaryl, CN, optionally substituted CQ<sup>a</sup>Q<sup>b</sup>aryl, optionally substituted CQ<sup>a</sup>Q<sup>b</sup>heteroaryl or COR<sup>7</sup>;

R<sup>7</sup> represents hydrogen, optionally substituted alkyl, optionally substituted heteroaryl or optionally substituted aryl;

R<sup>8</sup> and R<sup>9</sup> are independently selected from hydrogen, fluorine or alkyl, or R<sup>8</sup> and R<sup>9</sup> together with the carbon to which they are attached form a cycloalkyl ring, optionally containing up to one heteroatom selected from O, S, NH or N-alkyl;

wherein Q<sup>a</sup> and Q<sup>b</sup> are each independently selected from hydrogen, CH<sub>3</sub> and fluorine;

or a derivative thereof.

- 2. (Original) A compound according to claim 1 wherein the five membered ring comprising W, X and Y is pyrrole or pyrazole.
- 3. (Currently Amended) A compound according to claim 1 or claim 2 wherein  $R^1$  is  $CO_2H$ .
- 4. (Canceled).
- 5. (Currently Amended) A pharmaceutical composition comprising a compound according to any one of claims 1 to 4 or a pharmaceutically acceptable derivative thereof together with a pharmaceutical carrier and/or excipient.
- 6. 7. (Canceled).
- 8. (Currently Amended) A method of treating a human or animal subject suffering from a condition which is mediated by the action of PGE<sub>2</sub> at EP<sub>1</sub>

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receptors which comprises administering to said subject an effective amount of a compound according to any one of claims 1 to 4 or a pharmaceutically acceptable derivative thereof.

- 9. (Currently Amended) A method of treating a human or animal subject suffering from inflammatory pain, neuropathic pain or visceral pain which method comprises administering to said subject an effective amount of a compound according to any one of claims 1 to 4 or a pharmaceutically acceptable derivative thereof.
- 10. 11. (Canceled).
- 12. (New) The method of claim 8, wherein the subject is a human.
- 13. (New) The method of claim 9, wherein the subject is a human.
- 14. (New) A method of mediating EP<sub>1</sub> receptors, comprising the step of administering an effective amount of a compound according to claim 1 or a pharmaceutically acceptable derivative thereof.